

L4 ANSWER 13 OF 13 CAPLUS COPYRIGHT 2003 ACS on STN
 AN 1994:30773 CAPLUS
 DN 120:30773
 TI Oxadiazole derivatives having acetylcholinesterase-inhibitory
 and muscarinic receptor agonist activity
 IN Takasugi, Hisashi; Kuno, Atsushi; Ohkubo, Mitsuru
 PA Fujisawa Pharmaceutical Co., Ltd., Japan
 SO PCT Int. Appl., 149 pp.
 CODEN: PIXXD2

DT Patent
 LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9313083	A1	19930708	WO 1992-JP1658	19921218
	W: AU, CA, HU, JP, KR, RU, US			GB, GR, IE, IT, LU, MC, NL, PT, SE	
	RW: AT, BE, CH, DE, DK, ES, FR,			GB 1991-27533	19911231
				GB 1992-20904	19921005
	AU 9331714	A1	19930728	AU 1993-31714	19921218
				GB 1991-27533	19911231
				GB 1992-20904	19921005
				WO 1992-JP1658	19921218
	EP 619814	A1	19941019	EP 1993-900416	19921218
	R: AT, BE, CH, DE, DK, ES, FR,			GB, GR, IE, IT, LI, LU, NL, PT, SE	
				GB 1991-27533	19911231
				GB 1992-20904	19921005
				WO 1992-JP1658	19921218
	JP 07502529	T2	19950316	JP 1992-511547	19921218
				GB 1991-27533	19911231
				GB 1992-20904	19921005
				WO 1992-JP1658	19921218
	US 5622976	A	19970422	US 1994-244904	19940624
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Patel

<10/13/2003>

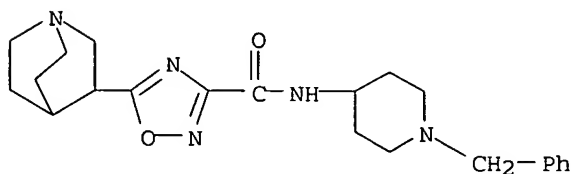
OS MARPAT 120:30773

IT 151097-86-8P 151097-87-9P 151307-60-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. and acetylcholinesterase inhibitory and muscarinic receptor
 agonist activity of)

RN 151097-86-8 CAPLUS

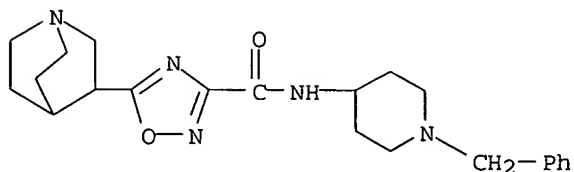
CN 1,2,4-Oxadiazole-3-carboxamide, 5-(1-azabicyclo[2.2.2]oct-3-yl)-N-[1-
 (phenylmethyl)-4-piperidiny]-, dihydrochloride (9CI) (CA INDEX NAME)



● 2 HCl

RN 151097-87-9 CAPLUS

CN 1,2,4-Oxadiazole-3-carboxamide, 5-(1-azabicyclo[2.2.2]oct-3-yl)-N-[1-
 (phenylmethyl)-4-piperidiny]- (9CI) (CA INDEX NAME)



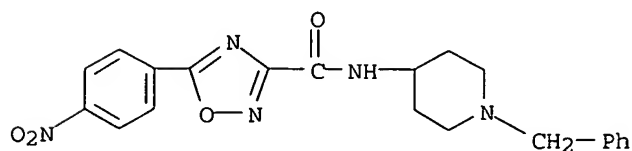
RN 151307-60-7 CAPLUS

CN 1,2,4-Oxadiazole-3-carboxamide, 5-(4-nitrophenyl)-N-[1-(phenylmethyl)-4-
 piperidiny]-, (2E)-2-butenedioate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 151307-59-4

CMF C21 H21 N5 O4



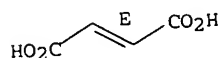
CM 2

10069215.2

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CRN -110-17-8
CMF C4 H4 O4

Double bond geometry as shown.



AB The title compds. R1QZXAM [A = direct bond, lower alkylene, lower alkynylene; M = (un)substituted heterocyclic group contg. 1 N atom(s); Q = oxadiazole-diyl; R1 = lower alkyl, (un)substituted heterocyclic group, (un)substituted aryl, (un)substituted arylalkyl, (un)substituted aralkenyl; X = direct bond, CONR4, R8CN; R4 = H, alkyl; R8 = HO, protected HO group, CO, NHCO; Z = direct bond, vinyl (sic)], useful for the treatment of central nervous system disorders (e.g., amnesia, Alzheimer's disease, vascular dementia, etc.) mode data, are prepd. Thus, 3-ethoxycarbonyl-5-(quinuclidin-3-yl)-1,2,4-oxadiazole and 1-benzyl-4-(2-aminoethyl)piperidine were heated together in soln. at 100.degree. for 2 h and treated with an ethanolic soln. of HCl, producing 5-(quinuclidin-3-yl)-3-[[2-(1-benzylpiperidin-4-yl)ethyl]carbamoyl]-1,2,4-oxadiazole dihydrochloride, m.p. 210.degree. (decompn.).